## Amendments to the Claims:

Please amend Claims 1-3 and 33, cancel Claim 3, and add new Claims 60-67. This listing of claims will replace all prior versions, and listings of claims in the application:

## **Listing of Claims:**

1

2

1 (currently amended): A method for identifying a therapeutic agent for use in 1 treating a constitutive androstane receptor (CAR)-mediated disorder or condition, wherein the 2 CAR-mediated disorder or condition is hypercholesterolemia, the method comprising: 3 identifying a candidate therapeutic agent by screening one or more compounds to 4 determine whether said compounds ean modulate comprise an agonist of a CAR-mediated 5 6 intermolecular interaction; administering the candidate therapeutic agent to a test mammal; and 7 determining whether the level of a cholesterol indicator is modulated decreased in 8 said test mammal in comparison to a test mammal in which the candidate therapeutic agent is not 9 administered. 10

2 (currently amended): The method of claim 1, wherein said candidate therapeutic agent is selected from the group consisting of 5ß-pregnan-3,20-dione, 1,4-bis[2-(3,5-

3 dichloropyridyloxy)]benzene (TCPOBOP),

, and mixtures thereof.

3 (canceled).

5

- 4 (previously presented): The method of claim 1, wherein the test mammal is a cholesterol-elevated mammal.
- 5 (original): The method of claim 4, wherein the test mammal has a disruption in both CAR alleles.
- 1 6 (original): The method of claim 1, wherein said cholesterol indicator is the level of serum cholesterol.
- 7 (original): The method of claim 1, wherein said cholesterol indicator is the level of a member selected from the group consisting of HDL cholesterol, LDL cholesterol, and VLDL cholesterol.
- 8 (original): The method of claim 1, wherein said cholesterol indicator is the mRNA level of a gene involved in the regulation of cholesterol levels.
- 9 (original): The method of claim 1, wherein said CAR-mediated intermolecular interaction is CAR-mediated gene expression.

## 10-32 (canceled)

| 1  | 33 (currently amended): A method for identifying a therapeutic agent for use in               |  |  |
|----|---|--|--|
| 2  | treating a constitutive androstane receptor (CAR)-mediated disorder or condition, wherein the |  |  |
| 3  | CAR-mediated disorder or condition is hypercholesterolemia, the method comprising:            |  |  |
| 4  | administering a compound to a CAR compromised mammal, wherein said CAR                        |  |  |
| 5  | compromised mammal comprises a mutation, disruption or insertion in at least one CAR allele   |  |  |
| 5  | that prevents the production of a functional CAR polypeptide; and                             |  |  |
| 7  | determining whether administration of the compound results in a change in                     |  |  |
| 8  | cholesterol level compared to a CAR compromised mammal to which the compound is not           |  |  |
| 9  | administered.   |  |  |
| 1  | 34 (original): The method of claim 33, wherein the method further comprises                   |  |  |
| 2  | administering the compound to a CAR non-compromised mammal and comparing the effect on        |  |  |
| 3  | the cholesterol level indicator of administering the compound to that of administering the    |  |  |
| 4  | compound to the CAR compromised mammal.   |  |  |
| 1  | 35 (original): The method of claim 33, wherein said cholesterol level indicator is            |  |  |
| 2  | the level of serum cholesterol.   |  |  |
| 1  | 36 (original): The method of claim 33, wherein said cholesterol level indicator is            |  |  |
| 2  | the level of a member selected from the group consisting of HDL cholesterol, LDL cholesterol, |  |  |
| 3  | and VLDL cholesterol.   |  |  |
| 1. | 37 (original): The method of claim 33, wherein said cholesterol level indicator is            |  |  |
| 2  | the mRNA level of a gene involved in the regulation of cholesterol levels.                    |  |  |
| 1  | 38 (original): The method of claim 33, wherein said CAR compromised mamma                     |  |  |
| 2  | is a mammal having a disruption in both CAR alleles.  |  |  |

| 1  | 39 (original): The method of claim 38, wherein said CAR compromised mammal  |  |  |
|----|---|--|--|
| 2  | is a mouse.   |  |  |
| 1  | 40 (original): The method of claim 38, wherein said disruption occurs in the  |  |  |
| 2  | coding region for the DNA binding domain of CAR.  |  |  |
| 1  | 41 (original): The method of claim 38, wherein said disruption in a CAR allele  |  |  |
| 2  | comprises an insertion at codons for amino acid positions from about amino acid 21 to about                                       |  |  |
| 3  | amino acid 86 of CARß.  |  |  |
|    | 42-59 (canceled)  |  |  |
| 1  | 60. (new) The method of claim 1, wherein said CAR-mediated intermolecular   |  |  |
| 2  | interaction comprises CAR binding to a ligand for CAR.  |  |  |
| 1  | 61. (new) A method for identifying a therapeutic agent for use in treating a  |  |  |
| 2  | constitutive androstane receptor (CAR)-mediated disorder or condition, wherein the CAR-   |  |  |
| 3  | mediated disorder or condition is hypocholesterolemia, the method comprising:   |  |  |
| 4  | identifying a candidate therapeutic agent by screening one or more compounds to   |  |  |
| 5  | determine whether said compounds comprise at least one of an antagonist or an inverse agonist                                     |  |  |
| 6  | of a CAR-mediated intermolecular interaction;   |  |  |
| 7  | administering the candidate therapeutic agent to a test mammal; and   |  |  |
| 8  | determining whether the level of a cholesterol indicator is increased in said test  |  |  |
| 9  | mammal in comparison to a test mammal in which the candidate therapeutic agent is not   |  |  |
| 10 | administered.   |  |  |
| 1  | 62. (new) The method of claim 61, wherein said candidate therapeutic agent is   |  |  |
| 2  | selected from the group consisting of $5\alpha$ -androst-16-en- $3\alpha$ -ol, $5\alpha$ -androstane- $3\alpha$ -ol, androstenol- |  |  |
| 3  | 3-acetate, $5\alpha$ -androstan- $3\alpha$ -ol-acetate, androstenol, and mixtures thereof.  |  |  |
|    |   |  |  |

| 1 | 63.   | (new) The method of claim 61, wherein said cholesterol indicator is the |  |
|---|---|---|--|
| 2 | level of serum cholesterol.   |   |  |
| 1 | , 64.   | (new) The method of claim 61, wherein said cholesterol indicator is the |  |
| 2 | level of a member selected from the group consisting of HDL cholesterol, LDL cholesterol, and |   |  |
| 3 | VLDL cholesterol.   |   |  |
| 1 | 65.   | (new) The method of claim 61, wherein said cholesterol indicator is the |  |
| 2 | mRNA level of a gene involved in the regulation of cholesterol levels.                        |   |  |
| 1 | 66.   | (new) The method of claim 61, wherein said CAR-mediated                 |  |
| 2 | intermolecular interaction is CAR-mediated gene expression.                                   |   |  |
| 1 | 67.   | (new) The method of claim 61, wherein said CAR-mediated                 |  |
| 2 | intermolecular interaction comprises CAR binding to a ligand for CAR.                         |   |  |